Exhibit I

Deposition of Terri-Lee Nataline
December 14, 2009

UNITED STATES DISTRICT COURT

FOR THE SOUTHERN DISTRICT OF WEST VIRGINIA

CHARLESTON DIVISION

- - -

IN RE: DIGITEK® PRODUCTS LIABILITY LITIGATION : MDL NO. 1968

(This document relates to all cases.)

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CONFIDENTIAL - SUBJECT TO FURTHER CONFIDENTIALITY REVIEW

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New York, New York
Monday, December 14, 2009

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Videotaped Deposition of TERRI-LEE NATALINE, ESQUIRE, held at Harris Beach PLLC, 100 Wall Street, 24th Floor, on the above date, beginning at 9:34 a.m., before Kimberly A. Overwise, a Certified Realtime Reporter and Notary Public.

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manufacturing practices regulations that the FDA puts out, are they applicable to all the drugs that Actavis manufactures?

A Yes.

Q Are there any differences between the standards applicable to Digitek and those GMP standards applicable to the other drugs made by the company?

A No.

Q What's the purpose of GMPs?

MR. ANDERTON: Objection.

You may answer.

THE WITNESS: It's to ensure good manufacturing procedures. It's to make sure that, you know, the proper procedures are followed and the product is made in the appropriate manner.

BY MR. BLIZZARD:

Q Okay. Does it relate to safety?

MR. ANDERTON: Objection.

You may answer.

THE WITNESS: It depends -- it depends on -- potentially, yes.